

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 13, 2003, 12:46:48 ; Search time 1416 Seconds
(without alignments)
2166.823 Million cell updates/sec

Title: US-09-880-253B-5

Perfect score: 75

Sequence: 1 agacuccagcccgaccgc.....acaccuccucugagaccg 75

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*

- 1: gb_ba.*
- 2: gb_htg.*
- 3: gb_in.*
- 4: gb_om.*
- 5: gb_ov.*
- 6: gb_pat.*
- 7: gb_ph.*
- 8: gb_pl.*
- 9: gb_pr.*
- 10: gb_ro.*
- 11: gb_sts.*
- 12: gb_sy.*
- 13: gb_un.*
- 14: gb_vi.*
- 15: em_ba.*
- 16: em_fun.*
- 17: em_hum.*
- 18: em_in.*
- 19: em_mu.*
- 20: em_om.*
- 21: em_or.*
- 22: em_ov.*
- 23: em_pat.*
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- 31: em_htg_inv.*
- 32: em_htg_other.*
- 33: em_htg_mus.*
- 34: em_htg_pln.*
- 35: em_htg_rod.*
- 36: em_htg_nam.*
- 37: em_htg_vrt.*
- 38: em_sy.*
- 39: em_htgo_hum.*
- 40: em_htgo_mus.*
- 41: em_htgo_other.*

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	75	100.0	3600	6	BD141660	BD141660 Use of Gl
2	75	100.0	3600	6	BD141661	BD141661 Use of Gl
3	75	100.0	3600	6	BD141662	BD141662 Use of Gl
4	75	100.0	3600	6	BD141663	BD141663 Use of Gl
5	75	100.0	3600	9	HSGLI	X07384 Human mRNA
6	48	64.0	1492	9	AF026306	AF026306 Homo sapi
7	48	64.0	166987	2	AC063917	AC063917 Homo sapi
8	48	64.0	172136	9	AC022506	AC022506 Homo sapi
9	48	64.0	185688	2	AC018805	AC018805 Homo sapi
10	40	53.3	201672	2	AC122965	AC122965 Rattus no
11	40	53.3	207478	2	AC108599	AC108599 Rattus no
12	40	53.3	218231	2	AC114111	AC114111 Rattus no
13	39.6	52.8	3587	6	AX676836	AX676836 Sequence
14	39.6	52.8	3587	9	BC013000	BC013000 Homo sapi
15	36.8	49.1	796	10	AF189287	AF189287 Mus muscu
16	36.8	49.1	206936	2	AC114678	AC114678 Mus muscu
17	34.2	45.6	8513	6	AX251122	AX251122 Sequence
18	34.2	45.6	8513	6	AX277897	AX277897 Sequence
19	34.2	45.6	8513	6	AX323574	AX323574 Sequence
20	34.2	45.6	8513	6	AX344687	AX344687 Sequence
21	31.4	41.9	158811	2	AC013503	AC013503 Homo sapi
22	31.4	41.9	175773	9	AC027128	AC027128 Homo sapi
23	31.4	41.9	192973	9	AC023471	AC023471 Homo sapi
24	31.4	41.9	222542	9	AC023379	AC023379 Homo sapi
25	30	40.0	22970	9	HSL247F6	Z68279 Human DNA S
26	30	40.0	174318	9	CNS01RGQ	AL159141 Human chr
27	30	40.0	193371	2	AC098298	AC098298 Rattus no
28	30	40.0	209157	9	CNS01DW4	AL136332 Human chr
29	30	40.0	219949	2	AC113819	AC113819 Rattus no
30	29.8	39.7	2080	6	AX302031	AX302031 Sequence
31	29.8	39.7	114979	2	AC020798	AC020798 Mus muscu
32	29.8	39.7	139150	9	AC138655	AC138655 Homo sapi
33	29.8	39.7	217304	9	AC009600	AC009600 Homo sapi
34	29.8	39.7	265234	2	AC102562	AC102562 Mus muscu
35	29.2	38.9	1620	6	AX440610	AX440610 Sequence
36	29.2	38.9	110000	2	AC091338	AC091338 Rattus no
37	29.2	38.9	220722	2	AC116257	AC116257 Rattus no
38	29.2	38.9	221507	9	HS407F11	AL022329 Human DNA
39	29.2	38.9	233675	2	AC132796	AC132796 Rattus no
40	29.2	38.9	234542	2	AC127639	AC127639 Rattus no
41	29.2	38.9	236139	2	AC125724	AC125724 Rattus no
42	29.2	38.9	237090	2	AC094943	AC094943 Rattus no
43	29.2	38.9	266201	2	AC091336	AC091336 Rattus no
44	29	38.7	385	9	AF005644	AF005644 Pan trogl
45	29	38.7	385	9	AF005645	AF005645 Pan panis

ALIGNMENTS

RESULT 1	BD141660	BD141660	3600 bp	DNA	linear	PAT 18-SEP-2002
LOCUS	BD141660	Use of Glil gene.				
DEFINITION	BD141660	BD141660				
ACCESSION	BD141660.1	GI:23236605				
VERSION	WO 0211752-A/10.					
KEYWORDS	Homo sapiens (human)					
SOURCE	Homo sapiens					
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;					
REFERENCE	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.					
AUTHORS	1 (bases 1 to 3600)					
TITLE	Hikichi, Y.					
JOURNAL	Use of Glil gene					
	Patent: WO 0211752-A 10 14-FEB-2002;					

	TAKEDA CHEMICAL INDUSTRIES LTD,YUICHI HIKICHI
COMMENT	OS Homo sapiens (human)
PN	WO 0211752-A/10
PD	14-FEB-2002
PF	03-AUG-2001 WO 2001JP006688
PR	04-AUG-2000 JP OOP 242767
PI	YUICHI HIKICHI
PC	A61K38/17,A61K48/00,A61K45/00,A61P19/00,A61P19/02,A61P19/10,
PC	C12Q1/02,
PC	C12Q1/68,G01N33/15,G01N33/50//C07K14/47,C12N15/12 CC
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Best Local Similarity	89.3%; Pred.No. 2.7e-11;
Matches	67; Conservative 8; Mismatches 0; Indels 0; Gaps 0
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Dd	4 AGACTCCAGCGCTGGACCGCGCATCCGAGCCCCAGCGCCCAGACAGAGTGTCCCCAACC 63 : : : : : : : :
QY	61 CUCCUCUGAGACGCC 75
Dd	::: :
Dd	64 CTCCTCGAGACGCC 78
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LOCUS	BD141661
DEFINITION	Use of Gli1 gene.
ACCESSION	BD141661
VERSION	BD141661.1 GI:23236606
KEYWORDS	WO 0211752-A/11.
SOURCE	Homo sapiens (human)
ORGANISM	Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 3600) Hikichi,Y. Use of Gli1 gene Patent: WO 0211752-A 11 14-FEB-2002; TAKEDA CHEMICAL INDUSTRIES LTD,YUICHI HIKICHI
COMMENT	OS Homo sapiens (human)
PN	WO 0211752-A/11
PD	14-FEB-2002
PF	03-AUG-2001 WO 2001JP006688
PR	04-AUG-2000 JP OOP 242767
PI	YUICHI HIKICHI
PC	A61K38/17,A61K48/00,A61K45/00,A61P19/00,A61P19/02,A61P19/10,
PC	C12Q1/02,
PC	C12Q1/68,G01N33/15,G01N33/50//C07K14/47,C12N15/12 CC
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	/mol_type="genomic DNA"
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VTKHRGDGPLPRAFSISTVEPKREREGGPIRESRLTVPEGAMKQPSGPAQSSCSS
DHSPAGSAANTSGVMTGNAGGSTDLSLDEGPCIAGTGLSTLRLENLRLDQLHQ
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PQYLQSGPYTQPPPDYLPSEPRPCLDFDSTHSTGQLKAQLVCNVVQSQELLWEGG
REDAPAEPSYQSPKFLGGSQVSPRAKAPVNTYGPFGNLPNHKSGSYPTPSPCHE
NFVVGANRASHRAAAPRLPLPTCYGPKLVGGTNPSCGHPVEVGRUGGPPALYPPE
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Best Local Similarity 89.3%; Pred. No. 2.7e-11;
Matches 67; Conservative 8; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCUGGACCGGCAUCCCGAGCCCGCCAGACAGAGUGUCCCCACACC 60
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 4 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGCCAGACAGAGTGTCCCCACACC 63

QY 61 CUCCUCUGAGACGCC 75
|:|:|:|:|:|:|:|
Db 64 CTCCTCTGAGACGCC 78

RESULT 5
HSGLI
LOCUS HSGLI 3600 bp mRNA linear PRI 12-SEP-1993
DEFINITION Human mRNA for GLI protein.
ACCESSION X07384
VERSION X07384.1 GI:31767
KEYWORDS GLI protein; zinc finger protein.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 3600)
AUTHORS Kinzler,K.W., Ruppert,J.M., Bigner,S.H. and Vogelstein,B.
TITLE The GLI gene is a member of the Kruppel family of zinc finger
proteins
JOURNAL Nature 332 (6162), 371-374 (1988)
MEDLINE 88175051
PUBMED 2832761
REFERENCE 2 (bases 1 to 3600)
AUTHORS Kinzler,K.W.
TITLE Direct Submission
JOURNAL Submitted (03-MAY-1988)
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Location/Qualifiers
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785 a 1162 c 948 g 705 t
BASE COUNT 785 a 1162 c 948 g 705 t
ORIGIN
Query Match 100.0%; Score 75; DB 6; Length 3600;
Best Local Similarity 89.3%; Pred. No. 2.7e-11;
Matches 67; Conservative 8; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCUGGACCGGCAUCCCGAGCCCGCCAGACAGAGUGUCCCCACACC 60
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 4 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGCCAGACAGAGTGTCCCCACACC 63

QY 61 CUCCUCUGAGACGCC 75
|:|:|:|:|:|:|:|
Db 64 CTCCTCTGAGACGCC 78

RESULT 6
AF026306
LOCUS AF026306 1492 bp DNA linear PRI 23-SEP-1999
DEFINITION Homo sapiens zinc finger transcription factor GLI (GLI) gene,
5'UTR, partial sequence.
ACCESSION AF026306
VERSION AF026306.2 GI:5919240
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1492)
AUTHORS Liu,C.Z., Yang,J.T., Yoon,J.W., Villavicencio,E., Pfendler,K.,
Walterhouse,D. and Iannaccone,P.
TITLE Characterization of the promoter region and genomic organization of
GLI, a member of the Sonic hedgehog-Patched signaling pathway
JOURNAL Gene 209 (1-2), 1-11 (1998)
MEDLINE 98192509
PUBMED 9524201
REFERENCE 2 (bases 1 to 1492)
AUTHORS Liu,C.Z., Yang,J.T., Yoon,J.W., Walterhouse,D. and Iannaccone,P.
TITLE Direct Submission
JOURNAL Submitted (23-SEP-1997) Pediatrics, Northwestern University Medical
School/Children's Memorial Hospital, 2300 Children's Plaza,
Chicago, IL 60614, USA
REFERENCE 3 (bases 1 to 1492)
AUTHORS Liu,C.Z., Yang,J.T., Yoon,J.W., Walterhouse,D. and Iannaccone,P.
TITLE Direct Submission
JOURNAL Submitted (23-SEP-1999) Pediatrics, Northwestern University Medical
School/Children's Memorial Hospital, 2300 Children's Plaza,
Chicago, IL 60614, USA
REMARK
COMMENT On Sep 23, 1999 this sequence version replaced gi:3004846.

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* 76220 89425: contig of 13206 bp in length
* 89426 89525: gap of unknown length
* 89526 109142: contig of 19617 bp in length
* 109143 109242: gap of unknown length
* 109243 130358: contig of 21116 bp in length
* 130359 130458: gap of unknown length
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Matches 45; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
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Db 154715 AGACTCCAGCCCTGGACCGCGCATCCGAGCCCGAGCCCGCCAGACAGAG 154668
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LOCUS
DEFINITION Homo sapiens 12 BAC RP11-181L23 (Roswell Park Cancer Institute
Human BAC Library) complete sequence.
AC022506
AC022506.38 GI:24418014
HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 172136)
AUTHORS Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,
Alsbrooks,S.L., Amaratunge,H.C., Are,J.R., Ayele,M., Banks,T.,
Barbaria,J., Benton,J., Bimage,K., Blankenburg,K., Bonnin,D.,
Bouck,J., Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P.,
Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C.,
Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D.,
Chen,G., Chen,R., Chen,Z., Chiu,D., Chowdhry,I., Christopoulos,C.,
Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R.,
Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A.,
Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H.,
Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Durbin,K.J.,
Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Emerling,S.,
Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P.,
Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N.,
Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S.,
Hamilton,K., Han,J., Harris,C., Harris,K., Hart,M., Havlak,P.,
Hawes,A., Hernandez,J., Hernandez,O., Hodgson,A., Hogues,M.,
Holloway,C., Hollins,B., Homsí,F., Howard,S., Huber,J., Hulyk,S.,
Hume,J., Ioshikhes,I., Jackson,L.E., Jacobson,B., Jia,Y.,
Johnson,R., Jolivet,S., Joudah,S., Karlsson,E., Kelly,S., Khan,U.,
King,L., Korvah,J., Kovar,C., Kratovic,J., Kureshi,A., Landry,N.,
Leal,B., Lee,E., Lewis,L.C., Lewis,L., Li,J., Li,Z., Lichtarge,O.,
Lieu,C., Liu,J., Liu,W., Loulseged,H., Lozado,R.J., Lu,X.,
Lucier,A., Lucier,R., Luna,R., Ma,J., Maheshwari,M., Mapua,P.,
Marondel,I., Martin,R., Martindale,A., Martinez,E., Massey,E.,
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Mohabbat,K., Montgomery,K.T., Morgan,M., Morris,S., Moser,M.,
Neal,D., Nelson,D., Newton,J., Newton,N., Nguyen,A., Nguyen,N.,
Nguyen,N., Nickerson,E., Nwokenkwu,S., Oguh,M., Okwuonu,G.,
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Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y.,
Rives,M., Rojas,A., Rojibokan,I., Rolfe,M., Ruiz,S., Savery,G.,
Scherer,S., Scott,G., Shen,H., Shim,C., Shooshtari,N., Sisson,I.,

Sodergren,E., Sonaik,T., Sparks,A., Stanley,H., Stone,H.,
Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H.,
Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S.,
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Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S.,
Williams,G., Williamson,A., Wleczyk,R., Wooden,S., Worley,K.,
Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Kucherlapati,R.,
Weinstock,G. and Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 172136)
Worley,K.C.
Direct Submission
Submitted (05-FEB-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 172136)
Worley,K.C.
Direct Submission
Submitted (11-SEP-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 172136)
Worley,K.C.
Direct Submission
Submitted (29-OCT-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
5 (bases 1 to 172136)
Worley,K.C.
Direct Submission
Submitted (26-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
6 (bases 1 to 172136)
Worley,K.C.
Direct Submission
Submitted (15-MAR-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Oct 29, 2002 this sequence version replaced gi:22779458.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email
gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
entire insert of this clone. Overlapping regions of clones are only
sequenced and submitted once, so the sequence for the remainder of
the insert may be found in the record for the adjacent clones.
Overlapping clones are noted at the beginning and end of the
Features listing.

ANNOTATION OF FEATURES:

STGs are identified using ePCR (Genome Res. 7:541-550) searches
of a local database that includes entries from dbSTS, GDB, and
local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green,
unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST
(Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the
EST and cDNA sequences. Genes demonstrate at least two exons
flanked by consensus splice sites that maintained sequence
continuity across the splice junctions. Sequences that are not
identical matches are annotated as similar.

SEQUENCING READ COVERAGE:Sequencing is completed to a minimum
standard of double strand coverage with a minimum of 2 clones and 2
reads with no ambiguities or 2 chemistries with a minimum of 2
clones and 3 reads with no ambiguities. If the sequence quality for
a region does not meet this standard, it will be indicated in the
annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES:This sequence meets stringent quality
standards - estimated error rate less than 1 per 10,000 bases.

Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

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STS

2773. .3082

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repeat_region

4627. .4649
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repeat_region

complement(9542. .9847)
/rpt_family="AluSx"

repeat_region

9889. .10004
/rpt_family="FLAM_A"

repeat_region

10023. .11439
/rpt_family="L2"

repeat_region

complement(11572. .11882)
/rpt_family="AluSp"

repeat_region

complement(12363. .12663)
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repeat_region

complement(12988. .13081)
/rpt_family="MIR"

repeat_region

complement(13082. .13398)
/rpt_family="AluSx"

repeat_region

complement(13399. .13516)
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repeat_region

14379. .14410
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repeat_region

14874. .15037
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repeat_region

16935. .16970
/rpt_family="GC-rich"

repeat_region

complement(17292. .17654)
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repeat_region

17657. .17909
/rpt_family="AluSx"

repeat_region

17949. .18167
/rpt_family="AluSg/x"

repeat_region

18168. .18244
/rpt_family="AluSx"

repeat_region

18245. .19209
/rpt_family="LTR13"

repeat_region

19210. .19443
/rpt_family="AluSx"

repeat_region

complement(19480. .20171)
/rpt_family="L2"

repeat_region

20325. .20574
/rpt_family="AluSx"

STS

21607. .21773

repeat_region

/standard_name="RH103508"

repeat_region

22151. .22171
/rpt_family="(A)n"

repeat_region

23441. .23494
/rpt_family="(TG)n"

STS

23604. .23839

repeat_region

/standard_name="D12S1889"
complement(25085. .25168)
/rpt_family="MIR"

repeat_region complement(25816. .25988)
/rpt_family="FAM"
STS 28112. .28315
/standard_name="25471"
repeat_region complement(28275. .28323)
/rpt_family="MER66A"
repeat_region complement(29010. .29304)
/rpt_family="AluJb"
repeat_region complement(30472. .30773)

Query Match 64.0%; Score 48; DB 9; Length 172136;
Best Local Similarity 93.8%; Pred. No. 0.00048;
Matches 45; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCCGGACCGCGCAUCCCGAGCCCGCCAGACAGAG 48
Db 147932 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGCCAGACAGAG 147885

RESULT 9
AC018805/c

LOCUS AC018805 185688 bp DNA linear HTG 07-JUL-2000
DEFINITION Homo sapiens chromosome 12 clone RP11-564P5, WORKING DRAFT
SEQUENCE, 29 unordered pieces.
AC018805
VERSION AC018805.4 GI:8568931
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 185688)
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 185688)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (03-JAN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

COMMENT On Jun 16, 2000 this sequence version replaced gi:6855245.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc/index.shtml>

----- Project Information -----
Center project name: H_NH0564P05
----- Summary Statistics -----

Sequencing vector: M13; 87%
Sequencing vector: plasmid; 13%
Chemistry: Dye-primer ET; 87% of reads
Chemistry: Dye-terminator Big Dye; 13% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 172728 bases at least Q40
Consensus quality: 176882 bases at least Q30
Consensus quality: 179092 bases at least Q20
Insert size: 217000; agarose-fp
Insert size: 182888; sum-of-contigs
Quality coverage: 3.63 in Q20 bases; agarose-fp
Quality coverage: 4.26 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 29 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 1241: contig of 1241 bp in length
* 1242 1341: gap of unknown length

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

1 (bases 1 to 201672)

Muzny,D.Marie., Metzker,M.Lee., Abramzon,S., Adams,C., Alder,J., Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D., Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H., Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F., Biswalo,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M., Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E., Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A., Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J., Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L., Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D., Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K., Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K., Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G., Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P., Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M., Gregeorge,M., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W., Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K., Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J., Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hognes,M., Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A., Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A., Karpathy,S., Kelly,S., Khan,Z., King,L., Kovar,C., Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J., Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J., Loreshuhewa,L., Loulseghe,H., Lozado,R.J., Lu,X., Ma,J., Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A., Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E., Mawhiney,S., McLeod,M.P., McNeill,T.Z., Meenen,E., Molasavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S., Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L., Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S., Nwaokemeleh,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K., Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkoch,C., Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.-L., Puazo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R., Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F., Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J., Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., Shen,H., Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D., Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J., Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C., Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K., Valas,R., Vera,V., Villasan,D., Waldron,L., Walker,B., Wang,J., Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F., Williams,G., Willson,R., Wleczyk,R., Wooden,H., Worley,K., Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V., Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von Niederhausen,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O., Weinstock,G. and Gibbs,R.A.

Direct Submission

Unpublished

2 (bases 1 to 201672)

Worley,K.C.

Direct Submission

Submitted (26-MAY-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 201672)

Rat Genome Sequencing Consortium.

Direct Submission

Submitted (19-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On Nov 19, 2002 this sequence version replaced gi:23907835. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold,

individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GYCT

Center clone name: CH230-349N19

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 175103 bases at least Q40

Consensus quality: 177389 bases at least Q30

Consensus quality: 178394 bases at least Q20

Estimated insert size: 180387; sum-of-contigs estimation

Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 2 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 16227: contig of 16227 bp in length

* 16228 16327: gap of unknown length

* 16328 201672: contig of 185345 bp in length.

FEATURES

source

1. 201672

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-349N19"

/complement(14639..15474)

/note="clone boundary"

clone_end:Sp5

site:

end sequence:BZ181092"

16328..17800

/note="wgs_contig"

BASE COUNT 47238 a 43520 c 42628 g 45829 t 22457 others

ORIGIN

Query Match

Best Local Similarity 53.3%; Score 40; DB 2; Length 201672;

Matches 40; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCCGGACCGCGCAUCCGAGCCGCGCCGACGACAGAG 48

Db 89400 AGTTCCAGCCCTGGACCGCATCCCGAGCCCTGCGCCCGACAGAG 89447

RESULT 11

AC108599/c

LOCUS

AC108599 Rattus norvegicus clone CH230-299C4, WORKING DRAFT SEQUENCE. 207478 bp DNA linear HTG 11-OCT-2002

DEFINITION

AC108599

VERSION

AC108599.4 GI:23603444

KEYWORDS

HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_FULLTOP.

SOURCE

Rattus norvegicus (Norway rat)

ORGANISM

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus.

KEYWORDS
SOURCE
ORGANISM

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.
Rattus norvegicus (Norway rat)

Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE
AUTHORS

1 (bases 1 to 218231)
Muzny,D.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gebregorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Haylak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hogue,M.,
Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpathy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensuhewa,L., Loulseged,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhiney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwaokenleh,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkoch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.-L.,
Puazo,M., Quiróz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wlarczyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

Unpublished
2 (bases 1 to 218231)
Worley,K.C.
Direct Submission
Submitted (07-MAR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

REFERENCE
AUTHORS
TITLE
JOURNAL

3 (bases 1 to 218231)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (13-MAY-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

COMMENT

On May 13, 2003 this sequence version replaced gi:23268081.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas

(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu

----- Project Information
Center project name: GLMH
Center clone name: CH230-23P10

----- Summary Statistics
Assembly program: Atlas 3.0;

Consensus quality: 185672 bases at least Q40

Consensus quality: 190509 bases at least Q30

Consensus quality: 193980 bases at least Q20

Estimated insert size: 194985; sum-of-contigs estimation

Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length

* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 5 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 121740: contig of 121740 bp in length

* 121741 121840: gap of unknown length

* 121841 195708: contig of 73868 bp in length

* 195709 195808: gap of unknown length

* 195809 214376: contig of 18568 bp in length

* 214377 214476: gap of unknown length

* 214477 216260: contig of 1784 bp in length

* 216261 216360: gap of unknown length

* 216361 218231: contig of 1871 bp in length.

FEATURES

Source

1. 218231

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-23P10"

890. 1591

/note="clone boundary

clone end:Sp6

site:ECORI

end_sequence:BH275857"

complement(17653..18480)

/note="clone boundary

clone end:T7

site:ECORI

end_sequence:BH275856"

27885..29132

/note="wgs_end_extension

clone end:T7"

32824..35023

/note="wgs_end_extension

clone end:T7"

36172..37544

/note="wgs_end_extension

clone end:T7"

202535..204207

/note="wgs_end_extension

clone_end:T7"

BASE COUNT 49557 a 48073 c 48428 g 50669 t 21504 others
ORIGIN

Query Match 53.3%; Score 40; DB 2; Length 218231;
Best Local Similarity 83.3%; Pred. No. 0.083;
Matches 40; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCCGGACCGCGCAUCCCGAGCCCGAGCCCGCCAGACAGAG 48
|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Db 9123 AGTTCCAGCCCTGGACCAGCATCCCGAGCCCTGGCCCGCCGACAGAG 9170

RESULT 13
AX676836
LOCUS AX676836 3587 bp DNA linear PAT 27-MAR-2003
DEFINITION Sequence 29 from Patent WO02103028.
ACCESSION AX676836
VERSION AX676836.1 GI:29334445
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Baranova,A.V., Yankovsky,N.K., Kozlov,A.P., Lobashev,A.V. and Krukovskaya,L.L.
TITLE In silico screening for phenotype-associated expressed sequences
JOURNAL Patent: WO 02103028-A 29 27-DEC-2002;
Biomedical Center (RU)

FEATURES
source Location/Qualifiers
1..3587
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
BASE COUNT 798 a 1143 c 944 g 702 t
ORIGIN

Query Match 52.8%; Score 39.6; DB 6; Length 3587;
Best Local Similarity 80.4%; Pred. No. 0.26;
Matches 37; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 30 GCCCAGGCCCGCCAGACAGAGUGUCCCGACACCCUCCUGAGAGCGCC 75
|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Db 2 GCACGAGGCCCGCCAGACAGAGTGTCGCCACACCCCTCTCTGAGAGCGCC 47

RESULT 14
BC013000
LOCUS BC013000 3587 bp mRNA linear PRI 22-AUG-2001
DEFINITION Homo sapiens, Similar to glioma-associated oncogene homolog (zinc finger protein), clone MGC:3590 IMAGE:3531657, mRNA, complete cds.
ACCESSION BC013000
VERSION BC013000.1 GI:15278120
KEYWORDS MGC.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Strausberg,R.
TITLE Direct Submission
JOURNAL Submitted (20-AUG-2001) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA
REMARK NIH-MGC Project URL: http://mgc.nci.nih.gov
COMMENT Contact: MGC help desk
Email: cgabbs-remail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbiology.org

contact: amadan@systemsbiology.org
Anup Madan, Rachel Dickhoff, Jessica Fahey, Stephanie Ford, Julia Greene, Mark Kettman and Anuradha Madan

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Series: IRAL Plate: 11 Row: h Column: 7
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 4885278.

FEATURES
Location/Qualifiers
1..3587
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="MGC:3590 IMAGE:3531657"
/tissue type="Muscle, rhabdomyosarcoma"
/clone_lib="NIH_MGC_17"
/lab_host="DH10B-R"
/note="Vector: pOTB7"
48..3368
/codon_start=1
/product="Similar to glioma-associated oncogene homolog (zinc finger protein)"
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BASE COUNT 798 a 1143 c 944 g 702 t
ORIGIN

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Best Local Similarity 80.4%; Pred. No. 0.26;
Matches 37; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

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|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Db 2 GCACGAGGCCCGCCAGACAGAGTGTCGCCACACCCCTCTCTGAGAGCGCC 47

RESULT 15
AF189287
LOCUS AF189287 796 bp DNA linear ROD 28-OCT-1999
DEFINITION Mus musculus zinc finger transcription factor GLI (Gli) gene, promoter and 5'UTR, partial sequence.
ACCESSION AF189287
VERSION AF189287.1 GI:6137222

KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 796)
AUTHORS Walterhouse,D., Ahmed,M., Slusarski,D., Kalamaras,J., Boucher,D., Holmgren,R. and Iannaccone,P.
TITLE gli, a zinc finger transcription factor and oncogene, is expressed

during normal mouse development
Dev. Dyn. 196 (2), 91-102 (1993)
MEDLINE 93372381
PUBMED 8364225
REFERENCE 2 (bases 1 to 796)
AUTHORS Liu,C.Z., Yang,J.T., Yoon,J.W., Villavicencio,E., Pfendler,K.,
Walterhouse,D. and Iannaccone,P.
TITLE Characterization of the promoter region and genomic organization of
GLI, a member of the Sonic hedgehog-Patched signaling pathway
JOURNAL Gene 209 (1-2), 1-11 (1998)
MEDLINE 98192509
PUBMED 9524201
REFERENCE 3 (bases 1 to 796)
AUTHORS Liu,C.Z., Yang,J.T., Yoon,J.W., Villavicencio,E., Pfendler,K.,
Walterhouse,D. and Iannaccone,P.
TITLE Direct Submission
JOURNAL Submitted (23-SEP-1999) Pediatrics, Children's Memorial Institute
for Education and Research, 2430 N. Halsted St., CMIER C-503,
Chicago, IL 60614, USA
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 /db_xref="taxon:10090"
 /chromosome="10"
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 mRNA
 242..>796
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 242..>796
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 5'UTR
BASE COUNT 152 a 215 c 261 g 168 t
ORIGIN
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Best Local Similarity 79.2%; Pred. No. 2.2;
Matches 38; Conservative 3; Mismatches 7; Indels 0; Gaps 0;
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Search completed: November 13, 2003, 13:18:13
Job time : 1420 secs

GenCore version 5.1.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 13, 2003, 12:45:23 ; Search time 254 Seconds
(without alignments)
797.078 Million cell updates/sec

Title: US-09-880-253B-5
Perfect score: 75
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Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 2552756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	75	100.0	75	ABK30106	Human GLI gene, Hg
2	75	100.0	3600	AAD12302	Human Cubitus inte
3	75	100.0	3600	AAL45542	Human Gli1 coding
4	75	100.0	3600	AAL45543	Human Gli1 coding
5	75	100.0	3600	AAL45544	Human Gli1 coding
6	75	100.0	3600	AAL45545	Human Gli1 coding
7	75	100.0	3600	ABK30501	Human glioma-assoc
8	73.4	97.9	75	ABK30157	Human Hgamma-UTR p

9	71.8	95.7	75	24	ABK30158	Huma Hgamma-UTR pl
10	70.2	93.6	75	24	ABK30159	Human Hbgamma-UTR
11	50.2	66.9	219	24	ABK30105	Human GLI gene, Hb
12	48.6	64.8	219	24	ABK30154	Human Hbeta-UTR pl
13	48	64.0	581	24	ABK30136	Human GLI UTR part
14	48	64.0	1492	24	ABK30508	Human glioma-assoc
15	48	64.0	4620	24	ABK30161	Human GLI1 genomic
16	47	62.7	219	24	ABK30155	Huma Hbeta-UTR plu
17	47	62.7	219	24	ABK30156	Human Hbeta-UTR pl
18	45.4	60.5	74	24	ABK30104	Mouse GLI gene, Mg
19	43.8	58.4	74	24	ABK30151	Mouse Mgamma-UTR p
20	42.2	56.3	74	24	ABK30152	Mouse Mgamma-UTR p
21	40.6	54.1	74	24	ABK30153	Mouse Mgamma-UTR p
22	36.8	49.1	188	24	ABK30103	Mouse GLI gene, Mb
23	36.8	49.1	307	24	ABK30102	Mouse GLI gene, Ma
24	36.8	49.1	307	24	ABK30142	Mouse Malpha-UTR.
25	36.8	49.1	877	24	ABK30134	Mouse GLI UTR part
26	36.8	49.1	3707	24	ABK30160	Mouse GLI1 genomic
27	35.2	46.9	188	24	ABK30148	Mouse Mbeta-UTR pl
28	34.2	45.6	8513	22	AAS45355	Chemically pretrea
29	34.2	45.6	8513	22	AAS46368	Tumour suppressor
30	34.2	45.6	8513	24	ABN80095	Human chemically m
31	34.2	45.6	8513	24	ABK28188	DNA transcription
32	33.6	44.8	188	24	ABK30149	Mouse Mbeta-UTR pl
33	33.6	44.8	188	24	ABK30150	Mouse Mbeta-UTR pl
34	33.6	44.8	307	24	ABK30143	Mouse Malpha-UTR p
35	30.4	40.5	307	24	ABK30144	Mouse Malpha-UTR p
36	29.8	39.7	2080	24	AAD22007	Human transporters
37	29.6	39.5	770	21	AAC70773	Single nucleotide
38	29.6	39.5	770	21	AAC70782	Single nucleotide
39	29.6	39.5	770	21	AAC70785	Single nucleotide
40	29.6	39.5	770	21	AAC70791	Single nucleotide
41	29.2	38.9	1620	24	ABA99034	Human G protein-co
42	29	38.7	573	24	ABK30137	Human GLI UTR part
43	29	38.7	33654	22	AAF70259	Human dopamine rec
44	29	38.7	33654	25	ABT34216	Human dopamine rec
45	28.8	38.4	307	24	ABK30145	Mouse Malpha-UTR p

ALIGNMENTS

RESULT 1
ABK30106
ID ABK30106 standard; RNA; 75 BP.
XX
AC ABK30106;
XX 23-APR-2002 (first entry)
DT Human GLI gene, Hgamma-UTR mRNA.
XX
KW Human; mouse; gene therapy; pseudo-translation initiation site;
KW herbicide resistance; pesticide resistance; transgenic plant; ss.
XX Homo sapiens.
OS
XX WO200196569-A1.
PN
XX 20-DEC-2001.
PD
XX 13-JUN-2001; 2001WO-AU00697.
PF
XX 13-JUN-2000; 2000US-211159P.
PR
XX (UYQU) UNIV QUEENSLAND.
PA
XX Rothnagel JA, Wang X;
PI
XX WPI; 2002-098072/13.
DR
XX Modulating expression of genetic sequence, comprising ORF having
PT RTG/RUG corresponding to authentic translation site, involves

PT introducing/removing RTG/RUG triplets in nucleotide sequence upstream
PT of authentic site -
XX
PS Claim 51; Page 103; 147pp; English.
XX
CC The invention relates to a method of modulating expression of a genetic
CC sequence, comprising introducing, creating or deleting one or more
CC pseudo-translation initiation sites, in the nucleotide sequence of an
CC mRNA, 5' upstream of the authentic translation initiation site of an open
CC reading frame (ORF), or by introducing, creating or deleting Kozac
CC sequences genetically proximal to the pseudo-translation initiation
CC sites. The method is useful for modulating the expression of a target
CC genetic sequence. The method is useful for gene therapy applications and
CC for expressing traits (herbicide and pesticide resistance) at selective
CC levels in plants. The genetic constructs are useful for administration
CC to modulate the expression of an antigen. The method is also useful for
CC the generation of a genetically modified monocotyledon or dicotyledon
CC plants, and also for upregulating or downregulating the function of a
CC promoter. ABK30102-ABK30161 represent human and mouse GLI gene sequences
CC and PCR primers of the invention.
XX
SQ Sequence 75 BP; 15 A; 35 C; 17 G; 8 U; 0 other;

Query Match 100.0%; Score 75; DB 24; Length 75;
Best Local Similarity 100.0%; Pred. No. 1.4e-13;
Matches 75; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCCGGCGCAUCCCGAGCCGCGCCAGACAGAGUGUCCCCACACC 60
Db |||||
1 AGACUCCAGCCCGGCGCAUCCCGAGCCGCGCCAGACAGAGUGUCCCCACACC 60

QY 61 CUCCUCUGAGACGCC 75
Db |||||
61 CUCCUCUGAGACGCC 75

RESULT 2
AAD12302
ID AAD12302 standard; cDNA; 3600 BP.
XX
AC AAD12302;
XX
DT 16-OCT-2001 (first entry)
XX
DE Human Cubitus interruptus (Ci) homologue, GLI-1 cDNA.
XX
KW Human; transgenic non-human animal; Cubitus interruptus; Ci; GLI-1;
KW basal cell carcinoma; BCC model system; tumour; screening;
KW anti-cancer; trichoeptithelioma; cylindroma; trichoblastoma; ss.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
CDS 79..3399
FT /*tag= a
FT /product= "Human Ci homologue, GLI-1"
XX
PN WO200156376-A1.
XX
XX
PD 09-AUG-2001.
XX
PF 02-FEB-2001; 2001WO-SE00204.
XX
PR 03-FEB-2000; 2000SE-0000345.
XX
PA (KARO-) KAROLINSKA INNOVATIONS AB.
XX
PI Toftgard R;
XX
DR WPI; 2001-488828/53.
DR P-PSDB; AAB06644.
XX
PT Transgenic non-human animal useful as basal cell carcinoma model system

PT to identify anti-cancer drug candidates, overexpresses transgene
PT encoding GLI-1 protein which is a human homolog to Cubitus interruptus
PT -
XX
PS Claim 6; Page 25-26; 33pp; English.
XX
CC The present invention relates to a transgenic non-human animal
CC comprising a transgene containing a nucleic acid encoding a human
CC Cubitus interruptus (Ci) homologue protein, GLI-1. The transgenic
CC non-human animal is useful as basal cell carcinoma (BCC) model system
CC since it overexpresses GLI-1 which leads to development of tumours
CC resembling human BCC. Thus it is also useful for screening anti-cancer
CC drug candidates and evaluating whether it affects BCC,
CC trichoeptitheliomas, cylindromas and trichoblastomas. The present
CC sequence is a cDNA encoding human Ci homologue protein, GLI-1.
XX
SQ Sequence 3600 BP; 785 A; 1161 C; 949 G; 705 T; 0 other;

Query Match 100.0%; Score 75; DB 22; Length 3600;
Best Local Similarity 89.3%; Pred. No. 1.8e-13;
Matches 67; Conservative 8; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCCGGCGCAUCCCGAGCCGCGCCAGACAGAGUGUCCCCACACC 60
Db |||||
4 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCGCGCCAGACAGAGTGTCCCCACACC 63

QY 61 CUCCUCUGAGACGCC 75
Db |||||
64 CTCCTCTGAGACGCC 78

RESULT 3
AAL45542
ID AAL45542 standard; cDNA; 3600 BP.
XX
AC AAL45542;
XX
DT 11-JUN-2002 (first entry)
XX
DE Human Gli1 coding sequence SEQ ID NO: 12.
XX
KW Gli1; screening method; bone induction; cartilage induction;
KW orthopaedic disease; dental disease; osteoporosis; hyperosteoegenesis;
KW osteopathic; antiarthritic; vulnary; immunosuppressive; human;
KW hyperchondrogenesis; gene; ss.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
CDS 79..3399
FT /*tag= a
FT /product= "Gli1"
XX
PN WO200211752-A1.
XX
PD 14-FEB-2002.
XX
PF 03-AUG-2001; 2001WO-JP06688.
XX
PR 04-AUG-2000; 2000JP-0242767.
XX
PA (TAKE) TAKEDA CHEM IND LTD.
XX
PI Hikichi Y;
XX
DR WPI; 2002-241709/29.
DR P-PSDB; AAO17109.
XX
PT Promotion of bone and cartilage formation using Gli1 protein or DNA
PT encoding it for treatment of skeletal disorders -
XX
PS Claim 6; Page 100-101; 154pp; Japanese.
XX


```
CC described in the exemplification of the invention.
XX
SQ Sequence 3600 BP; 786 A; 1161 C; 948 G; 705 T; 0 other;
    Query Match      100.0%; Score 75; DB 24; Length 3600;
    Best Local Similarity 89.3%; Pred. No. 1.8e-13;
    Matches 67; Conservative 8; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCCGGCGCAUCCCGAGCCCGCCAGCCAGAGUGUCCCCACACC 60
    |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Db 4 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGCCAGAGAGTGTCCCCACACC 63
    |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||

QY 61 CUCCUCUGAGACGCC 75
    |||:|||||:|||||
Db 64 CTCCTCTGAGACGCC 78

RESULT 6
AAL45545
ID AAL45545 standard; cDNA; 3600 BP.
XX
AC AAL45545;
XX
DT 11-JUN-2002 (first entry)
XX
DE Human Gli1 coding sequence SEQ ID NO: 18.
XX
KW Gli1; screening method; bone induction; cartilage induction;
KW orthopaedic disease; dental disease; osteoporosis; hyperosteo genesis;
KW osteopathic; antiarthritic; vulnery; immunosuppressive; human;
KW hyperchondrogenesis; gene; ss.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
CDS 79..3399
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FT /product= "Gli1"
FT /transl_except= (pos:2416..2599,aa:Trp)
FT /note= "there is an apparent 180 nucleotide
FT insertion between residues 780 and 781 of the Gli1
FT protein"
XX
PN WO200211752-A1.
XX
PD 14-FEB-2002.
XX
PF 03-AUG-2001; 2001WO-JP06688.
XX
PR 04-AUG-2000; 2000JP-0242767.
XX
PA (TAKE ) TAKEDA CHEM IND LTD.
XX
PI Hikichi Y;
XX
WPI; 2002-241709/29.
DR P-PSDB; AAO17112.
XX
PT Promotion of bone and cartilage formation using Gli1 protein or DNA
PT encoding it for treatment of skeletal disorders -
XX
PS Claim 6; Page 123-125; 154pp; Japanese.
XX
CC The present invention relates to agents for the promotion of bone and
CC cartilage formation which contain as the active component a Gli1 protein
CC or a DNA encoding a Gli1 protein. The agents can be used in the
CC prevention, treatment and diagnosis of bone and cartilage disorders
CC including bone fractures, joint deformation, osteoarthritis,
CC osteoporosis, cartilage damage, trauma, bone formation defects, cartilage
CC formation defects, bone defects, dental disease, hyperosteo genesis and
CC hyperchondrogenesis, and for use in cosmetic and therapeutic bone
CC transplantation. The present sequence is a human Gli1 coding sequence
CC described in the exemplification of the invention.

XX
SQ Sequence 3600 BP; 785 A; 1162 C; 948 G; 705 T; 0 other;
    Query Match      100.0%; Score 75; DB 24; Length 3600;
    Best Local Similarity 89.3%; Pred. No. 1.8e-13;
    Matches 67; Conservative 8; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCCGGCGCAUCCCGAGCCCGCCAGCCAGAGUGUCCCCACACC 60
    |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Db 4 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGCCAGAGAGTGTCCCCACACC 63
    |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||

QY 61 CUCCUCUGAGACGCC 75
    |||:|||||:|||||
Db 64 CTCCTCTGAGACGCC 78

RESULT 7
ABK30501
ID ABK30501 standard; DNA; 3600 BP.
XX
AC ABK30501;
XX
DT 23-APR-2002 (first entry)
XX
DE Human glioma-associated oncogene-1 DNA sequence.
XX
KW Human; glioma-associated oncogene-1 associated disease; infection;
KW inflammation; tumour formation; cytostatic; antiinflammatory;
KW gene; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
CDS 79..3399
FT /*tag= a
FT /product= "Glioma-associated oncogene-1 protein"
XX
PN US6329203-B1.
XX
PD 11-DEC-2001.
XX
PF 08-SEP-2000; 2000US-0657042.
XX
PR 08-SEP-2000; 2000US-0657042.
XX
PA (ISIS-) ISIS PHARM INC.
XX
PI Bennett CF, Wyatt J;
XX
WPI; 2002-138363/18.
DR P-PSDB; AAU12085.
XX
PT Novel antisense compounds targeted to nucleic acids encoding
PT glioma-associated oncogene-1, for modulating the gene expression and
PT treating diseases associated with expression of the oncogene in humans
PT -
XX
PS Example 13; Column 47-56; 43pp; English.
XX
CC The present invention relates to antisense compounds and methods for
CC modulating the expression of human glioma-associated oncogene-1. The
CC antisense compounds, particularly antisense oligonucleotides
CC (ABK30509-ABK30586), target and inhibit the expression of human
CC glioma-associated oncogene-1. The antisense compounds are useful for
CC inhibiting the expression of human glioma-associated oncogene-1 in human
CC cells or tissues and for treating an animal, particularly a human
CC suspected of having or being prone to a disease or condition associated
CC with expression of glioma-associated oncogene-1. The compounds are useful
CC for diagnostics, therapeutics and as research reagent,
CC e.g. prophylactically to prevent or delay infection, inflammation or
CC tumour formation. The antisense compounds are safely and effectively
CC administered to humans. The present sequence represents human
CC glioma-associated oncogene-1 DNA.
```


Db 61 CUCCUCUGAGACGCC 75

RESULT 10
ABK30159

ID ABK30159 standard; RNA; 75 BP.
XX
AC ABK30159;
XX
DT 23-APR-2002 (first entry)
XX
DE Human Hbgamma-UTR plus 3 ATG translation initiation site.
XX
KW Human; mouse; gene therapy; pseudo-translation initiation site;
KW herbicide resistance; pesticide resistance; transgenic plant; ss.
XX
OS Homo sapiens.
XX
PN WO200196569-A1.
XX
PD 20-DEC-2001.
XX
PF 13-JUN-2001; 2001WO-AU00697.
XX
PR 13-JUN-2000; 2000US-211159P.
XX
PA (UYQU) UNIV QUEENSLAND.
XX
PI Rothnagel JA, Wang X;
XX
DR WPI; 2002-098072/13.
XX
PS Modulating expression of genetic sequence, comprising ORF having
PT RTG/RUG corresponding to authentic translation site, involves
PT introducing/removing RTG/RUG triplets in nucleotide sequence upstream
PT of authentic site -
XX
XX Example 32; Page 76; 147pp; English.
XX
CC The invention relates to a method of modulating expression of a genetic
CC sequence, comprising introducing, creating or deleting one or more
CC pseudo-translation initiation sites, in the nucleotide sequence of an
CC mRNA, 5' upstream of the authentic translation initiation site of an open
CC reading frame (ORF), or by introducing, creating or deleting Kozac
CC sequences genetically proximal to the pseudo-translation initiation
CC sites. The method is useful for modulating the expression of a target
CC genetic sequence. The method is useful for gene therapy applications and
CC for expressing traits (herbicide and pesticide resistance) at selective
CC levels in plants. The genetic constructs are useful for administration
CC to modulate the expression of an antigen. The method is also useful for
CC the generation of a genetically modified monocotyledon or dicotyledon
CC plants, and also for upregulating or downregulating the function of a
CC promoter. ABK30102-ABK30161 represent human and mouse GLI gene sequences
CC and PCR primers of the invention.
XX
SQ Sequence 75 BP; 16 A; 32 C; 18 G; 9 U; 0 other;
Query Match 93.6%; Score 70.2; DB 24; Length 75;
Best Local Similarity 96.0%; Pred. No. 3.9e-12;
Matches 72; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCUGGACCGCGCAUCCCGAGCCCGCCAGACAGAGUGUCCCCACACC 60
Db 1 AGACUCCAGCCUGGACCGCGCAUCCCGAGCCCGCCAGACAGAGUGUCCCCACACC 60

QY 61 CUCCUCUGAGACGCC 75
Db 61 CUCCUCUGAGAUGCC 75

RESULT 11
ABK30105

ID ABK30105 standard; RNA; 219 BP.
XX
AC ABK30105;
XX
DT 23-APR-2002 (first entry)
XX
DE Human GLI gene, Hbeta-UTR mRNA.
XX
KW Human; mouse; gene therapy; pseudo-translation initiation site;
KW herbicide resistance; pesticide resistance; transgenic plant; ss.
XX
OS Homo sapiens.
XX
PN WO200196569-A1.
XX
PD 20-DEC-2001.
XX
PF 13-JUN-2001; 2001WO-AU00697.
XX
PR 13-JUN-2000; 2000US-211159P.
XX
PA (UYQU) UNIV QUEENSLAND.
XX
PI Rothnagel JA, Wang X;
XX
DR WPI; 2002-098072/13.
XX
PS Modulating expression of genetic sequence, comprising ORF having
PT RTG/RUG corresponding to authentic translation site, involves
PT introducing/removing RTG/RUG triplets in nucleotide sequence upstream
PT of authentic site -
XX
XX Claim 50; Page 103; 147pp; English.
XX
CC The invention relates to a method of modulating expression of a genetic
CC sequence, comprising introducing, creating or deleting one or more
CC pseudo-translation initiation sites, in the nucleotide sequence of an
CC mRNA, 5' upstream of the authentic translation initiation site of an open
CC reading frame (ORF), or by introducing, creating or deleting Kozac
CC sequences genetically proximal to the pseudo-translation initiation
CC sites. The method is useful for modulating the expression of a target
CC genetic sequence. The method is useful for gene therapy applications and
CC for expressing traits (herbicide and pesticide resistance) at selective
CC levels in plants. The genetic constructs are useful for administration
CC to modulate the expression of an antigen. The method is also useful for
CC the generation of a genetically modified monocotyledon or dicotyledon
CC plants, and also for upregulating or downregulating the function of a
CC promoter. ABK30102-ABK30161 represent human and mouse GLI gene sequences
CC and PCR primers of the invention.
XX
SQ Sequence 219 BP; 44 A; 65 C; 57 G; 53 U; 0 other;
Query Match 66.9%; Score 50.2; DB 24; Length 219;
Best Local Similarity 81.7%; Pred. No. 4.3e-06;
Matches 58; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 1 AGACUCCAGCCUGGACCGCGCAUCCCGAGCCCGCCAGACAGAGUGUCCCCACACC 60
Db 1 AGACUCCAGCCUGGACCGCGCAUCCCGAGCCCGCCAGACAGAGUGUCCCCACACC 60

QY 61 CUCCUCUGAGAGA 71
Db 61 UGUCUCACAGGA 71

RESULT 12
ABK30154

ID ABK30154 standard; RNA; 219 BP.
XX
AC ABK30154;
XX
DT 23-APR-2002 (first entry)
XX

DE Human Hbeta-UTR plus 1 ATG translation initiation site.
XX
KW Human; mouse; gene therapy; pseudo-translation initiation site;
KW herbicide resistance; pesticide resistance; transgenic plant; ss.
XX
OS Homo sapiens.
XX
PN WO200196569-A1.
XX
XX 20-DEC-2001.
PD
XX
PF 13-JUN-2001; 2001WO-AU00697.
XX
PR 13-JUN-2000; 2000US-211159P.
XX
XX (UYQU) UNIV QUEENSLAND.
PA
XX
PI Rothnagel JA, Wang X;
XX
DR WPI; 2002-098072/13.
XX
XX Modulating expression of genetic sequence, comprising ORF having
PT RTG/RUG corresponding to authentic translation site, involves
PT introducing/removing RTG/RUG triplets in nucleotide sequence upstream
PT of authentic site -
XX
PS Example 27; Page 74; 147pp; English.
XX
CC The invention relates to a method of modulating expression of a genetic
CC sequence, comprising introducing, creating or deleting one or more
CC pseudo-translation initiation sites, in the nucleotide sequence of an
CC mRNA, 5' upstream of the authentic translation initiation site of an open
CC reading frame (ORF), or by introducing, creating or deleting Kozac
CC sequences genetically proximal to the pseudo-translation initiation
CC sites. The method is useful for modulating the expression of a target
CC genetic sequence. The method is useful for gene therapy applications and
CC for expressing traits (herbicide and pesticide resistance) at selective
CC levels in plants. The genetic constructs are useful for administration
CC to modulate the expression of an antigen. The method is also useful for
CC the generation of a genetically modified monocotyledon or dicotyledon
CC plants, and also for upregulating or downregulating the function of a
CC promoter. ABK30102-ABK30161 represent human and mouse GLI gene sequences
CC and PCR primers of the invention.
XX
SQ Sequence 219 BP; 45 A; 64 C; 57 G; 53 U; 0 other;
Query Match 64.8%; Score 48.6; DB 24; Length 219;
Best Local Similarity 80.3%; Pred. No. 1.3e-05;
Matches 57; Conservative 0; Mismatches 14; Indels 0; Gaps 0;
QY 1 AGACUCCAGCCCGGCAUCCCGAGCCCGAGCCCGAGAGUCCCGACACC 60
Db 1 AGACUCCAGCCCGGCAUCCCGAGCCCGAGCCCGAGAGUCCCGACACC 60
QY 61 CUCCUCUGAGA 71
Db 61 UGUCUCAGGGA 71
RESULT 13
ABK30136
ID ABK30136 standard; DNA; 581 BP.
XX
AC ABK30136;
XX
DT 23-APR-2002 (first entry)
XX
DE Human GLI UTR partial genomic sequence #1.
XX
KW Human; mouse; gene therapy; pseudo-translation initiation site; gene;
KW herbicide resistance; pesticide resistance; transgenic plant; ds.
XX
OS Homo sapiens.

XX WO200196569-A1.
PN
XX 20-DEC-2001.
PD
XX
PF 13-JUN-2001; 2001WO-AU00697.
XX
PR 13-JUN-2000; 2000US-211159P.
XX
XX (UYQU) UNIV QUEENSLAND.
PA
XX
PI Rothnagel JA, Wang X;
XX
DR WPI; 2002-098072/13.
XX
XX Modulating expression of genetic sequence, comprising ORF having
PT RTG/RUG corresponding to authentic translation site, involves
PT introducing/removing RTG/RUG triplets in nucleotide sequence upstream
PT of authentic site -
XX
PS Example 7; Page 111; 147pp; English.
XX
CC The invention relates to a method of modulating expression of a genetic
CC sequence, comprising introducing, creating or deleting one or more
CC pseudo-translation initiation sites, in the nucleotide sequence of an
CC mRNA, 5' upstream of the authentic translation initiation site of an open
CC reading frame (ORF), or by introducing, creating or deleting Kozac
CC sequences genetically proximal to the pseudo-translation initiation
CC sites. The method is useful for modulating the expression of a target
CC genetic sequence. The method is useful for gene therapy applications and
CC for expressing traits (herbicide and pesticide resistance) at selective
CC levels in plants. The genetic constructs are useful for administration
CC to modulate the expression of an antigen. The method is also useful for
CC the generation of a genetically modified monocotyledon or dicotyledon
CC plants, and also for upregulating or downregulating the function of a
CC promoter. ABK30102-ABK30161 represent human and mouse GLI gene sequences
CC and PCR primers of the invention.
XX
SQ Sequence 581 BP; 114 A; 126 C; 196 G; 145 T; 0 other;
Query Match 64.0%; Score 48; DB 24; Length 581;
Best Local Similarity 93.8%; Pred. No. 2.1e-05;
Matches 45; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGACUCCAGCCCGGCAUCCCGAGCCCGAGCCCGAGAG 48
Db 1 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGAGCCCGAGAG 48
RESULT 14
ABK30508
ID ABK30508 standard; DNA; 1492 BP.
XX
AC ABK30508;
XX
DT 23-APR-2002 (first entry)
XX
DE Human glioma-associated oncogene-1 partial DNA sequence.
XX
KW Human; glioma-associated oncogene-1 associated disease; infection;
KW inflammation; tumour formation; cytostatic; antiinflammatory; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT CDS 1017..1067
FT /*tag= a
FT /partial
FT /product= "Partial peptide for human glioma-associated
FT oncogene-1"
FT /note= "This sequence lacks both start and stop codons"
XX
PN US6329203-B1.

XX PD 11-DEC-2001.
XX XX
XX PF 08-SEP-2000; 2000US-0657042.
XX XX
XX PR 08-SEP-2000; 2000US-0657042.
XX XX
XX PA (ISIS-) ISIS PHARM INC.
XX PI Bennett CF, Wyatt J;
XX XX
XX DR WPI; 2002-138363/18.
XX DR P-PSDB; AAU12086.
XX XX
XX PT Novel antisense compounds targeted to nucleic acids encoding
XX PT glioma-associated oncogene-1, for modulating the gene expression and
XX PT treating diseases associated with expression of the oncogene in humans
XX PT
XX XX
XX PS Example 15; Column 57-60; 43pp; English.
XX XX
XX CC The present invention relates to antisense compounds and methods for
XX CC modulating the expression of human glioma-associated oncogene-1. The
XX CC antisense compounds, particularly antisense oligonucleotides
XX CC (ABK30509-ABK30586), target and inhibit the expression of human
XX CC glioma-associated oncogene-1. The antisense compounds are useful for
XX CC inhibiting the expression of human glioma-associated oncogene-1 in human
XX CC cells or tissues and for treating an animal, particularly a human
XX CC suspected of having or being prone to a disease or condition associated
XX CC with expression of glioma-associated oncogene-1. The compounds are useful
XX CC for diagnostics, therapeutics and as research reagent,
XX CC e.g. prophylactically to prevent or delay infection, inflammation or
XX CC tumour formation. The antisense compounds are safely and effectively
XX CC administered to humans. The present sequence represents a partial
XX CC DNA sequence for human glioma-associated oncogene-1.
XX XX
XX SQ Sequence 1492 BP; 293 A; 415 C; 471 G; 313 T; 0 other;

Query Match 64.0%; Score 48; DB 24; Length 1492;
Best Local Similarity 93.8%; Pred. No. 2.2e-05;
Matches 45; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGACUCCAGCCUUGGACCGCGCAUCCCGAGCCCGCCAGACAGAG 48
Db 1020 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGCCAGACAGAG 1067

RESULT 15
ABK30161
ID ABK30161 standard; DNA; 4620 BP.
XX XX
XX AC ABK30161;
XX XX
XX DT 23-APR-2002 (first entry)
XX XX
XX DE Human GLI1 genomic sequence.
XX XX
XX KW Human; mouse; gene therapy; pseudo-translation initiation site; gene;
XX KW herbicide resistance; pesticide resistance; transgenic plant; ds.
XX OS Homo sapiens.
XX XX
XX PN WO200196569-A1.
XX XX
XX PD 20-DEC-2001.
XX XX
XX PF 13-JUN-2001; 2001WO-AU00697.
XX XX
XX PR 13-JUN-2000; 2000US-211159P.
XX XX
XX PA (UYQU) UNIV QUEENSLAND.
XX XX
XX PI Rothnagel JA, Wang X;

XX WPI; 2002-098072/13.
XX DR
XX XX
XX PT Modulating expression of genetic sequence, comprising ORF having
XX PT RTG/RUG corresponding to authentic translation site, involves
XX PT introducing/removing RTG/RUG triplets in nucleotide sequence upstream
XX PT of authentic site -
XX XX
XX PS Example 7; Page 120-122; 147pp; English.
XX XX
XX CC The invention relates to a method of modulating expression of a genetic
XX CC sequence, comprising introducing, creating or deleting one or more
XX CC pseudo-translation initiation sites, in the nucleotide sequence of an
XX CC mRNA, 5' upstream of the authentic translation initiation site of an open
XX CC reading frame (ORF), or by introducing, creating or deleting Kozac
XX CC sequences genetically proximal to the pseudo-translation initiation
XX CC sites. The method is useful for modulating the expression of a target
XX CC genetic sequence. The method is useful for gene therapy applications and
XX CC for expressing traits (herbicide and pesticide resistance) at selective
XX CC levels in plants. The genetic constructs are useful for administration
XX CC to modulate the expression of an antigen. The method is also useful for
XX CC the generation of a genetically modified monocotyledon or dicotyledon
XX CC plants, and also for upregulating or downregulating the function of a
XX CC promoter. ABK30102-ABK30161 represent human and mouse GLI gene sequences
XX CC and PCR primers of the invention.
XX XX
XX SQ Sequence 4620 BP; 916 A; 1354 C; 1368 G; 982 T; 0 other;

Query Match 64.0%; Score 48; DB 24; Length 4620;
Best Local Similarity 93.8%; Pred. No. 2.4e-05;
Matches 45; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGACUCCAGCCUUGGACCGCGCAUCCCGAGCCCGCCAGACAGAG 48
Db 1034 AGACTCCAGCCCTGGACCGCGCATCCCGAGCCCGCCAGACAGAG 1081

Search completed: November 13, 2003, 12:54:20
Job time : 254 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 13, 2003, 12:47:43 ; Search time 2051 Seconds
(without alignments)
888.755 Million cell updates/sec

Title: US-09-880-253B-5
Perfect score: 75
Sequence: 1 agacuccagcccgaccgc.....acaccuccucgagagcc 75

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

EST:*

- 1: em_estba:*
- 2: em_esthum:*
- 3: em_estin:*
- 4: em_estmu:*
- 5: em_estov:*
- 6: em_estpl:*
- 7: em_estro:*
- 8: em_htc:*
- 9: gb_est1:*
- 10: gb_est2:*
- 11: gb_htc:*
- 12: gb_est3:*
- 13: gb_est4:*
- 14: gb_est5:*
- 15: em_estfun:*
- 16: em_estom:*
- 17: em_gss_hum:*
- 18: em_gss_inv:*
- 19: em_gss_pln:*
- 20: em_gss_vrt:*
- 21: em_gss_fun:*
- 22: em_gss_mam:*
- 23: em_gss_mus:*
- 24: em_gss_pro:*
- 25: em_gss_rod:*
- 26: em_gss_phg:*
- 27: em_gss_vrl:*
- 28: gb_gss1:*
- 29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	39	52.0	767	12	BI114524
2	31	41.3	1008	13	BQ669921
C 3	30.6	40.8	364	14	R91407
C 4	30.4	40.5	812	13	BU424864

C 5	29.8	39.7	574	12	BI824472
C 6	29.8	39.7	575	12	BI547637
C 7	29.8	39.7	641	12	BI517462
C 8	29.8	39.7	669	12	BI757775
C 9	29.8	39.7	715	12	BI755911
C 10	29.8	39.7	1080	13	BX342864
C 11	29.8	39.7	1201	13	BX397064
C 12	29.6	39.5	407	12	BI724568
C 13	29.6	39.5	516	12	BM003088
C 14	29.4	39.2	563	29	BZ639961
C 15	29.4	39.2	867	10	BF616342
C 16	29.4	39.2	1000	13	BX342523
C 17	29.2	38.9	456	13	BX118143
C 18	29	38.7	390	9	AI123001
C 19	29	38.7	393	10	BG027933
C 20	29	38.7	745	12	BG827135
C 21	29	38.7	780	12	BI195997
C 22	28.8	38.4	265	10	BB585305
C 23	28.8	38.4	461	12	BM795944
C 24	28.8	38.4	463	10	BE394194
C 25	28.8	38.4	500	13	BU584167
C 26	28.8	38.4	511	9	AI692466
C 27	28.8	38.4	530	9	AW798307
C 28	28.8	38.4	576	12	BM700772
C 29	28.8	38.4	582	13	BU584165
C 30	28.8	38.4	666	10	BE394361
C 31	28.8	38.4	708	13	BX112900
C 32	28.8	38.4	730	12	BM044240
C 33	28.8	38.4	767	10	BG750003
C 34	28.8	38.4	877	13	BQ720868
C 35	28.8	38.4	916	12	BM019566
C 36	28.8	38.4	938	12	BI093058
C 37	28.8	38.4	943	13	BQ945928
C 38	28.8	38.4	1034	11	BC013276
C 39	28.6	38.1	320	9	AA024106
C 40	28.4	37.9	848	29	AG138174
C 41	28.4	37.9	1040	12	BM477912
C 42	28.2	37.6	389	12	BI261436
C 43	28.2	37.6	561	14	CA334682
C 44	28.2	37.6	601	12	BM041953
C 45	28.2	37.6	968	13	BQ681918

ALIGNMENTS

RESULT 1
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LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

BI114524
602862031F1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:5021359 5',
mRNA sequence.
BI114524
BI114524.1 GI:14565425
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 767)
NIH-MGC <http://mgs.nci.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
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LOCUS BI547637 575 bp mRNA linear EST 05-SEP-2001
DEFINITION 603191773F1 NIH_MGC_95 Homo sapiens cDNA clone IMAGE:5263002 5', mRNA sequence.
ACCESSION BI547637 GI:15434949
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 575)
NIH-MGC <http://mgc.nci.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki Toshiyuki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
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Location/Qualifiers
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/clone="IMAGE:5263002"
/tissue_type="hippocampus"
/lab_host="DH10B"
/clone_lib="NIH_MGC_95"
/note="Organ: brain; Vector: pBluescriptR (modified pBluescript KS+); Site_1: BamHI; Site_2: SalI-XhoI (gtcgag); Oligo-dT primed using primer 5'-TTTTTTT-TTTT-TTN-3', size-selected for average insert size 2.5 kb and normalized to ROT 5. This is a primary library enriched for full-length clones and constructed using the Cap-trapper method (Carninci, in preparation). Library constructed by M. Brownstein (NIH/NHGRI, National Institutes of Health). Note: this is a NIH_MGC Library."
BASE COUNT 68 a 222 c 230 g 55 t
ORIGIN
Query Match 39.7%; Score 29.8; DB 12; Length 575;
Best Local Similarity 58.9%; Pred. No. 2.7e+02;
Matches 43; Conservative 3; Mismatches 27; Indels 0; Gaps 0;
QY 2 GACUCCAGCCCGGCAUCCGAGCCCGCCAGACAGAGUGUCCCAACCC 61
Db 490 GCGGCCTTCGAGAGCCCTCGTCGCTTCCAGAGCCCGGACAGAGGGCGCCTCCGCC 431
QY 62 UCCUCUGAGACGC 74
Db 430 TCCTCCCCGGCGC 418
RESULT 7
BI517462/c
LOCUS BI517462 641 bp mRNA linear EST 29-AUG-2001
DEFINITION 603041736F1 NIH_MGC_116 Homo sapiens cDNA clone IMAGE:5162937 5', mRNA sequence.
ACCESSION BI517462 GI:15342254
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 641)

AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Plate: LLAM1404 row: h column: 10
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Location/Qualifiers
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/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5162937"
/lab_host="DH10B"
/clone_lib="NIH_MGC_116"
/note="Organ: pooled colon, kidney, stomach; Vector: pCMV-SPORT6; Site_1: NotI; Site_2: EcoRV (destroyed); RNA source anonymous pool of 3 colons, age 26 yo male, 49 yo female, 71 yo male colon; 46 yo male kidney, and pool of 2 stomachs, 62 yo male and 70 yo female. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.4 kb, insert size range 1-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 023. Note: this is a NIH_MGC Library."
BASE COUNT 76 a 246 c 256 g 63 t
ORIGIN
Query Match 39.7%; Score 29.8; DB 12; Length 641;
Best Local Similarity 58.9%; Pred. No. 2.7e+02;
Matches 43; Conservative 3; Mismatches 27; Indels 0; Gaps 0;
QY 2 GACUCCAGCCCGGCAUCCGAGCCCGCCAGACAGAGUGUCCCAACCC 61
Db 500 GCGGCCTTCGAGAGCCCTCGTCGCTTCCAGAGCCCGGACAGAGGGCGCCTCCGCC 441
QY 62 UCCUCUGAGACGC 74
Db 440 TCCTCCCCGGCGC 428
RESULT 8
BI757775/c
LOCUS BI757775 669 bp mRNA linear EST 25-SEP-2001
DEFINITION 603029745F1 NIH_MGC_114 Homo sapiens cDNA clone IMAGE:5200210 5', mRNA sequence.
ACCESSION BI757775 GI:15749353
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 669)
NIH-MGC <http://mgc.nci.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:

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http://image.llnl.gov
Plate: LLAM11501 row: i column: 11
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Location/Qualifiers
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/mol_type="mrna"
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/clone="IMAGE:5200210"
/lab_host="DH10B"
/clone_lib="NIH_MGC_114"
/note="Organ: brain; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: EcoRV (destroyed); RNA source anonymous pool of 6
male brains, age range 23-27 yo. Library is oligo-dT
primed and directionally cloned (EcoRV site is destroyed
upon cloning). Average insert size 1.5 kb, insert size
range 1-3 kb. Library is normalized and enriched for
full-length clones and was constructed by C. Gruber
(Invitrogen). Research Genetics tracking code 019. Note:
this is a NIH MGC Library."
86 a 258 c 256 g 69 t
BASE COUNT
ORIGIN

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Query Match		39.7%;	Score 29.8;	DB 12;	Length 669;
Best Local Similarity		58.9%;	Pred. No. 2.8e+02;		
Matches 43; Conservative		3;	Mismatches 27;	Indels 0;	Gaps 0;
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Dd	503	GCCGCCTTCGCAGAGCCCTCCTGTGGCCTTCCCGAGGCCCGGACAGAGGGGGCCTCGGC	444		
QY	62	UCCUCUGAGACGC	74		
Dd	443	TCCTCCCCGGCGC	431		

RESULT 9	BI755911/c	BI755911	715 bp	mRNA	linear	EST 25-SEP-2001					
LOCUS	603030127F1	NIH_MGC_114	Homo sapiens	cdna	clone	IMAGE:5200494 5',					
DEFINITION	mRNA sequence.										
ACCESSION	BI755911										
VERSION	BI755911.1	GI:15747489									
KEYWORDS	EST.										
SOURCE	Homo sapiens	(human)									
ORGANISM	Homo sapiens										
	Eukaryota;	Metazoa;	Chordata;	Craniata;	Vertebrata;	Euteleostomi;					
	Mammalia;	Eutheria;	Primates;	Catarrhini;	Hominidae;	Homo.					
1	(bases 1 to 715)										
NIH-MGC	http://mgc.nci.nih.gov/ .										
National Institutes of Health,	Mammalian Gene Collection (MGC)										
Unpublished											
Contact:	Robert Strausberg, Ph.D.										
Email:	cgapbs-r@mail.nih.gov										
Tissue Procurement:	Life Technologies, Inc.										
cdna Library Preparation:	Life Technologies, Inc.										
cdna Library Arrayed by:	The I.M.A.G.E. Consortium (LLNL)										
DNA Sequencing by:	Incyte Genomics, Inc.										
Clone distribution:	MGC clone distribution information can be										
found through the I.M.A.G.E. Consortium/LLNL at:	http://image.llnl.gov										
Plate:	LLAM11502	row: e	column: 07								
High quality sequence stop:	628.										
Location/Qualifiers											
1.	.715										
source											

```

FEATURES
source
    Location/Qualifiers
        1. .715
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="IMAGE:5200494"
            /lab_host="DH10B"
            /clone_lib="NIH_MGC_114"
            /note="Organ: brain; Vector: pCMV-SPORT6; Site_1: NotI;
            Site_2: EcoRV (destroyed); RNA source anonymous pool of 6

```

male brains, age range 23-27 yo. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.5 kb, insert size range 1-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 019. Note: this is a NIH MGCCLibrary."

```

BASE COUNT 123 a 256 c 267 g 69 t
ORIGIN

Query Match 39.7%; Score 29.8; DB 12; Length 715;
Best Local Similarity 58.9%; Pred. No. 2.8e+02;
Matches 43; Conservative 3; Mismatches 27; Indels 0; Gaps 0;

QY 2 GACUCCAGCCCUCCGACCGCGCAUCCCGAGCCCGCCAGACAGAGUGUCCCGACACCC 61
    ||||| ||||| : ||||| ||||| ||||| ||||| ||||| |||||
Db 505 GCGGCCTTCGACAGAGCCCCCTCGTCGCCTTCCACAGAGCCCGACAGAGGGGGCCTTCGCC 446

QY 62 UCCUCUGAGACGC 74
    :||:| |||||
Db 445 TCCTCCCCGGCGC 433

```

RESULT 10	
BX342864/c	
LOCUS	BX342864 1080 bp mRNA linear EST 02-MAY-2003
DEFINITION	BX342864 Homo sapiens B CELLS (RAMOS CELL LINE) COT 25-NORMALIZED Homo sapiens cDNA clone CS0DL007YC01 5-PRIME, mRNA sequence.
ACCESSION	BX342864
VERSION	BX342864.1 GI:30340126
KEYWORDS	EST.
SOURCE	Homo sapiens (human)
ORGANISM	Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 1080) Li,W.B., Gruber,C., Jessee,J. and Polayes,D. Full-length cDNA libraries and normalization Unpublished Contact: Genoscope
REFERENCE	
AUTHORS	
TITLE	
JOURNAL	
COMMENT	

```

Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr
Library was constructed by Life Technologies, a division of
Invitrogen. This sequence belongs to sequence cluster 1515.r For
more information about this cluster, see
http://www.genoscope.cns.fr/
cgi-bin/cluster.cgi?seq=CS0DL007AB01QP1&cluster=1515.r. Contact :
Feng Liang Email : fliang@lifetech.com URL :
http://fulllength.invitrogen.com/ InvitroGen Corporation 1600
Faraday Avenue Genoscope sequence ID : CS0DL007AB01QP1.
Location/Qualifiers
1. .1080
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CS0DL007YC01"
/cell_type="B CELLS (RAMOS CELL LINE) COT 25-NORMALIZED"
/cell_line="RAMOS CELL LINE"
/clone_lib="Homo sapiens B CELLS (RAMOS CELL LINE) COT
25-NORMALIZED"
/note="1st strand cDNA was primed with a NotI-oligo(dT)
primer. Five prime end enriched, double-strand cDNA was
digested with Not I and cloned into the Not I and EcoR V
sites of the pCMVSPORT 6 vector. Library was normalized."
131 a 402 c 386 q 132 t 29 others
BASE COUNT

```

```

BASE COUNT      131 a    402 c    386 g    132 t    29 others
ORIGIN

Query Match      39.7%;      Score 29.8;      DB 13;      Length 1080;
Best Local Similarity 58.9%;      Pred. No. 3e+02;
Matches 43; Conservative 3; Mismatches 27; Indels 0; Gaps 0;

QY      2 GACUCCAGCCCUUGGACCGGGCAUCCCTGAGCCCGCCAGCGCCCGAGACAGAGUGUCCCCACACCC 61

```


SK(-) CDNA phagemids. These steps were performed in the TJ
Close laboratory at the University of California,
Riverside (Choi, Close, Fenton). Phagemids were plated and
picked at the Clemson University Genomics Institute (CUGI)
(Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA
preparations, DNA sequencing and sequence analysis were
performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates
, Rambo, Main). The sequence has been trimmed to remove
vector sequence and contains a minimum of 100 bases of
phred value 20 or above. For more details on library
preparation and sequence analysis see
<http://www.genome.clemson.edu/projects/barley>. To order
this clone see <http://www.genome.clemson.edu/orders> Also
see Close TJ, Wing R, Kleinhofs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(<http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html>)"

BASE COUNT 153 a 123 c 408 g 183 t
ORIGIN

Query Match 39.2%; Score 29.4; DB 10; Length 867;
Best Local Similarity 61.9%; Pred. No. 3.7e+02;
Matches 39; Conservative 3; Mismatches 21; Indels 0; Gaps 0;

Qy 4 CUCCAGCCUUGGACCGCGCAUCCCGAGCCCGAGCCGAGAGAGUGUCCGACACCCUC 63
Db 832 CGCGGCCCCCGACCGATCAACCGGACCCCTGCCCGACCCACCTAACCCGCCACCTC 773

Qy 64 CUC 66
Db 772 CTC 770

Search completed: November 13, 2003, 13:52:38
Job time : 2055 secs